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☐ 1: Science. 1986 May 16;232(4752):854-8.

Science

Links

Cloning of a cDNA for a T cell-specific serine protease from a cytotoxic T lymphocyte.

Gershensfeld HK, Weissman IL.

A new serine protease was encoded by a clone isolated from a murine cytotoxic T-lymphocyte complementary DNA library by an RNA-hybridization competition protocol. Complementary transcripts were detected in cytotoxic T lymphocytes, spleen cells from nude mice, a rat natural killer cell leukemia, and in two of eight T-helper clones (both cytotoxic), but not in normal mouse kidney, liver, spleen, or thymus, nor in several tested T- and B-cell tumors. T-cell activation with concanavalin A plus interleukin-2 induced spleen cells to express this gene with kinetics correlating with the acquisition of cytolytic capacity. The nucleotide sequence of this gene encoded an amino acid sequence of approximately 25,700 daltons, with 25 to 35 percent identity to members of the serine protease family. The active site "charge-relay" residues (His57, Asp102, and Ser195 of the chymotrypsin numbering system) are conserved, as well as the trypsin-specific Asp (position 189 in trypsin). A Southern blot analysis indicated that this gene is conserved in humans, mouse, and chicken. This serine protease may have a role in lymphocyte lysis and a "lytic cascade."

PMID: 2422755 [PubMed - indexed for MEDLINE]

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A T cell- and natural killer cell-specific, trypsin-like serine protease. Implications of a lytic cascade. [Science. 1986]

Cloning and chromosomal assignment of a human cDNA encoding a T cell- and natural killer cell-specific trypsin-like serine protease. [Proc Natl Acad Sci U S A. 1988]

Characterization of a novel, human cytotoxic lymphocyte-specific serine protease cDNA. [Proc Natl Acad Sci U S A. 1990]

Novel serine proteases encoded by two cytotoxic T lymphocyte-specific genes. [Science. 1986]

Expression and utilization of chymotrypsin-like but not trypsin-like serine protease enzymes by nonspecific T killer cells activated by anti-CD3 monoclonal antibodies. [Cell. 1992]

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